

WANDERER DOCUMENTATION

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1. AIM

Wanderer is a very simple and intuitive web tool allowing real time access and visualization of gene expression and DNA methylation profiles obtained from the TCGA Research Network (<http://cancergenome.nih.gov/>) using gene targeted queries. Wanderer is addressed to a broad variety of experimentalists and clinicians without deep bioinformatics skills.

2. ACCESS AND CITATION

Wanderer may be accessed at <http://www.maplab.cat/wanderer>

If you find this software useful please consider citing our paper: (in preparation).

3. USING WANDERER

3.1. Enter a gene name (BRCA1 is used as default) or the equivalent Ensembl gene ID and press the Refresh button to update the output. Modifying the rest of options will automatically update the graphs.

3.2. Choose a Dataset (default dataset is Breast Invasive Carcinoma (BRCA)) and the Data type (450k Methylation array or Illumina HiSeq RNAseq) in the respective drop down menus.

3.3. Use the Zoom sliding bar to adjust the displayed chromosomal region or tick the Specify a region box for a more precise selection. Only values in the range defined by the zoom slider will be accepted.

3.4. Customize the graphical display of the data using Plotting parameters. Graphs are automatically updated after any change.

3.5. Download graphs and tables containing the selected data or share the output by generating a permanent link.

4. DESCRIPTION OF THE DOWNLOADABLE FILES GENERATED BY WANDERER

IMPORTANT WARNING ON USING EXCEL AND OTHER SPREADSHEET SOFTWARE TO IMPORT GENOMIC DATA

It is known that Excel and other spreadsheet softwares may change gene names when importing text data (Zeeberg et al., 2004). A default feature will misidentify specific gene names (e.g.: SET1, AGO3) as dates. This conversion may be avoided by formatting the spreadsheet cells as Text before importing.

Wanderer generates several images and data files that may be downloaded as a compressed file. The files are named using the following nomenclature:

```
Wanderer_[gene]_[data type]_[dataset]_[date and time of the query].
```

4.1 DNA METHYLATION FILES

The predefined query for DNA methylation produces the following graphs and data files:

4.1.1. Profile plots displaying the beta values for each probe in two separate panels representing subset of normal (upper panel, blue marks) and tumor samples (bottom panel, red marks) of the selected dataset. Lines link different probes corresponding to the same sample. The gene location and direction is depicted with an arrow. CpG island located probes are colored in green.

File name examples:

Raster version: 1_Wanderer_BRCA1_methylation_brca_Apr_29_2015_at_170440_CEST.png

Vectorial format: 2_Wanderer_BRCA1_methylation_brca_Apr_29_2015_at_170440_CEST.pdf

4.1.2. Two tables consisting of a comma separated data matrix containing data for normal and tumor samples respectively. Table contents: First column: 450K Methylation array probe ID, Rest of the columns: DNA methylation beta value for each sample.

File name examples:

3_Wanderer_BRCA1_methylation_brca_Normal_Apr_29_2015_at_170440_CEST.csv

4_Wanderer_BRCA1_methylation_brca_Tumor_Apr_29_2015_at_170440_CEST.csv

4.1.3. Profile plots displaying the average methylation for all the normal (blue) and tumor samples (red) of the selected dataset. The gene location and strand is depicted with an arrow. CpG island located probes are colored in green. The CpGs showing statistical differences between normal and tumor are highlighted with an asterisk (Wilcoxon adjusted p-value < adjusted pval threshold selected).

File name examples:

Raster version: 5_Wanderer_BRCA1_Mean_methylation_brca_Apr_29_2015_at_170440_CEST.png

Vectorial format: 6_Wanderer_BRCA1_Mean_methylation_brca_Apr_29_2015_at_170440_CEST.pdf

4.1.4. A comma separated data matrix with the annotation for each of the informative probes and descriptive analysis of the DNA methylation data. The columns correspond to:

- probe, probe name
- chr, the chromosome
- cg_start, the genomic position the CpG starts at
- cg_end, the genomic position the CpG ends at
- percentgc, the GC content of the illumina 450k array probe
- probetype, the type of the illumina 450k array probe
- probestart, the genomic position the probe starts at
- probeend, the genomic position the probe ends at
- genestart, the genomic position the closest gene to the probe starts at
- geneend, the genomic position the closest gene to the probe ends at
- genestrand, the closest gene strand
- ENSEMBL_geneID, the closest gene id at ensembl
- genebiotype, the closest gene biotype (protein coding, retained intron...)
- genename, the closest gene symbol
- cpgstart, for those CpG are inside a CpG island, the coordinate this islands starts at
- cpgiend, for those CpG are inside a CpG island, the coordinate this islands ends at
- cpgiid, for those CpG are inside a CpG island, CpG island identifier
- Norm_nsamples, number of normal samples for this dataset in this release
- Norm_mean, mean of the beta values for normals
- Norm_sd, standard deviation of the beta values for normals
- Tum_nsamples, number of tumor samples for this dataset in this release
- Tum_mean, mean of the beta values for tumors
- Tum_sd, standard deviation of the beta values for tumors
- wilcox_stat, if there are enough samples, Wilcoxon Rank Sum Test W parameter (nonparametric comparison of normals vs tumors)
- pval, if there are enough samples, Wilcoxon Rank Sum Test p value (nonparametric comparison of normals vs tumors, low values indicates that differences are detected)
- adj.pval, if there are enough samples, Benjamini and Hochberg adjustment (False Discovery Rate) for multiple testing.

File name examples:

[7_Wanderer_BRCA1_methylation_brca_annotations_and_statistical_analysis_Apr_29_2015_at_170440_CEST.csv](#)

4.1.5. Plots showing the correlation between DNA methylation beta values and RNAseq expression summarized by gene for all the normal (blue) and tumor samples (red) of the selected dataset. The expression values are log2-transformed normalized RSEM values (Guo et al., 2013) and reflect the expression of the gene as a whole. Three methods to obtain the correlation can be choosed: Spearman's rho, Kendall's tau and Pearson's r.

File name examples:

Raster version: [8_Wanderer_BRCA1_correl_RNAseqGeneVSmethylation_brca_Apr_29_2015_at_170440_CEST.png](#)

Vectorial format:

[9_Wanderer_BRCA1_correl_RNAseqGeneVSmethylation_brca_Apr_29_2015_at_170440_CEST.pdf](#)

4.1.6. A comma separated data matrix with the available normal and tumors sample names (first column) and the log2-transformed RSEM values for your gene of interest (second column).

File name examples:

10_Wanderer_BRCA1_methylation_brca_Normal_RNAseqGENE_CommonWithMethylation_Apr_29_2015_at_170440_CEST.c

11_Wanderer_BRCA1_methylation_brca_Tumor_RNAseqGENE_CommonWithMethylation_Apr_29_2015_at_170440_CEST.cs

4.2 GENE EXPRESSION FILES

4.2.1. Profile plots displaying the log2-transformed RPKM values (Guo et al., 2013) for each exon in two separate panels for a subset of normal (upper panel, blue marks) and tumor samples (bottom panel, red marks) from the selected dataset. Lines link values corresponding to the same sample. The gene location and strand is depicted with an arrow.

File name examples:

Raster version: 1_Wanderer_BRCA1_expression_brca_Apr_29_2015_at_170440_CEST.png

Vectorial format: 2_Wanderer_BRCA1_expression_brca_Apr_29_2015_at_170440_CEST.pdf

4.2.2. Two tables consisting of a comma separated data matrix containing data for normal and tumor samples respectively. Table contents: First column: RNAseq exon ID, Rest of the columns: log2-transformed RPKM values for each sample.

File name examples:

3_Wanderer_BRCA1_expression_brca_Normal_Apr_29_2015_at_170440_CEST.csv

4_Wanderer_BRCA1_expression_brca_Tumor_Apr_29_2015_at_170440_CEST.csv

4.2.3. Profile plots of the average expression for all the normal (blue) and tumor samples (red) of the selected dataset. The gene location and strand is depicted with an arrow. The exons showing statistically significant differences between normal and tumor are highlighted with an asterisk (Wilcoxon adjusted p-value < adjusted pval threshold selected).

File name examples:

Raster version: 5_Wanderer_BRCA1_Mean_expression_brca_Apr_29_2015_at_170440_CEST.png

Vectorial format: 6_Wanderer_BRCA1_Mean_expression_brca_Apr_29_2015_at_170440_CEST.pdf

4.2.4. A comma separated data matrix with the annotation of your gene expression data, as well some descriptive analysis. The columns correspond to:

- exon, exon identifier according to the TCGA pipeline
- id, exon identifier according to Genome Browser exons track
- ENSEMBL_geneID, the gene identifier according to ENSEMBL (ENSG identifier)
- ENSEMBL_transcriptID, the transcript identifier according to ENSEMBL (ENST identifier)
- chr, the chromosome the exon is located at
- exon_start, the genomic coordinate the exon starts at
- exon_end, the genomic coordinate the exon ends at
- strand, the genomic strand of the exon's gene.
- genestart, the genomic start position of the exon's gene.
- geneend, the genomic end position of the exon's gene.
- genebiotype, exon's gene biotype (protein coding, retained intron...)

- `genename`, exon's gene symbol
- `rnaseqgeneid`, the TCGA exon's gene identifier
- `Norm_nsamples`, number of normal samples for this dataset in this release
- `Norm_mean`, mean of log2-transformed RPKM for normals
- `Norm_sd`, standard deviation of log2-transformed RPKM in normals
- `Tum_nsamples`, number of tumor samples for this dataset in this release
- `Tum_mean`, mean of log2-transformed RPKM for tumors
- `Tum_sd`, standard deviation of log2-transformed RPKM in tumors
- `wilcox_stat`, if there are enough samples, Wilcoxon Rank Sum Test W parameter (nonparametric comparison of normals vs tumors)
- `pval`, if there are enough samples, Wilcoxon Rank Sum Test p value (nonparametric comparison of normals vs tumors, low values indicates that differences are detected)
- `adj.pval`, if there are enough samples, Benjamini and Hochberg adjustment (False Discovery Rate) for multiple testing.

File name example:

7_Wanderer_BRCA1_expression_brca_annotations_and_statistical_analysis_Apr_29_2015_at_170440_CEST.csv

4.2.5. Boxplot and a stripchart graphs showing the expression values summarized by gene for all the normal (blue) and tumor samples (red) of the selected dataset. The expression values are log2-transformed normalized RSEM values (Guo et al., 2013) and reflect the expression of the gene as a whole.

File name examples:

Raster version: 8_Wanderer_BRCA1_boxplot_expression_brca_Apr_29_2015_at_170440_CEST.png

Vectorial format: 9_Wanderer_BRCA1_boxplot_expression_brca_Apr_29_2015_at_170440_CEST.pdf

4.2.6. A comma separated data matrix with the available normal and tumors sample names (first column) and the log2-transformed RSEM values for your gene of interest (second column).

File name examples:

10_Wanderer_BRCA1_expression_brca_Normal_RNAseqGENE_Apr_29_2015_at_170440_CEST.csv

11_Wanderer_BRCA1_expression_brca_Tumor_RNAseqGENE_Apr_29_2015_at_170440_CEST.csv

4.3 CLINICAL DATA

TCGA clinical data is provided as biotab files. Biotab files are an amenable, spreadsheet-friendly data sources that reflect clinical and biospecimen information for a set of patients. In this file, each column is a clinical element and each row, a TCGA participant.

We note that biotab files cover data unique to each TCGA participant (for instance, the patient's weight and height) and therefore are independent on whether methylation or expression was scrutinized. We offer the biotab files for all the participants available for a dataset (for instance, colon adenocarcinoma), regardless of whether they were explored for expression or methylation, nor in solid primary tumors or in adjacent normal.

File name example:

BRCA_Clinical__nationwidechildrens.org_clinical_patient_brca.txt

5. DATA AVAILABILITY

Wanderer uses a TCGA snapshot taken on July 2014. Data availability for normals (N) and tumors (T) is detailed below.

Description	Name	Meth_N	Meth_T	RNAseq_N	RNAseq_T
Adrenocortical carcinoma	ACC	0	80	0	79
Bladder Urothelial Carcinoma	BLCA	21	358	19	267
Brain Lower Grade Glioma	LGG	0	511	0	513
Breast invasive carcinoma	BRCA	98	743	113	1052
Cervical squamous cell carcinoma and endocervical adenocarcinoma	CESC	3	256	3	207
Colon adenocarcinoma	COAD	38	302	41	262
Esophageal carcinoma	ESCA	16	185	0	0
Glioblastoma multiforme	GBM	2	129	5	156
Head and Neck squamous cell carcinoma	HNSC	50	528	43	497
Kidney Chromophobe	KICH	0	66	25	66
Kidney renal clear cell carcinoma	KIRC	160	324	72	518
Kidney renal papillary cell carcinoma	KIRP	45	226	30	198
Liver hepatocellular carcinoma	LIHC	50	256	50	212
Lung adenocarcinoma	LUAD	32	463	58	488
Lung squamous cell carcinoma	LUSC	43	361	50	491
Lymphoid Neoplasm Diffuse Large B+AC0-cell Lymphoma	DLBC	0	48	0	28
Mesothelioma	MESO	0	37	0	36
Ovarian serous cystadenocarcinoma	OV	0	10	0	262
Pancreatic adenocarcinoma	PAAD	10	146	4	96
Pheochromocytoma and Paraganglioma	PCPG	3	179	3	178
Prostate adenocarcinoma	PRAD	49	340	52	374
Rectum adenocarcinoma	READ	7	98	9	91

Description	Name	Meth_N	Meth_T	RNAseq_N	RNAseq_T
Sarcoma	SARC	4	242	2	103
Skin Cutaneous Melanoma	SKCM	2	92	1	82
Stomach adenocarcinoma	STAD	2	339	0	0
Thyroid carcinoma	THCA	56	507	59	498
Uterine Carcinosarcoma	UCS	0	57	0	57
Uterine Corpus Endometrial Carcinoma	UCEC	46	438	24	158
Uveal Melanoma	UVM	0	80	0	0

Wanderer offers external links to [Genome Browser](#), [Regulome Explorer](#) and [cBioPortal](#). We note that some of the user's queries might involve a gene or a dataset that was not covered in either Regulome Explorer or cBioPortal (i.e. mesothelioma). We note that the TCGA participants used in the analysis might differ between Wanderer and these others Web resources, as they used different data snapshots from TCGA. Moreover, Regulome Explorer datasets might contain tumor-only data (without the normal adjacent).

Description	Name	Regulome Explorer	cBioPortal
Adrenocortical carcinoma	ACC	acc_2015_03_31	acc_tcga
Bladder Urothelial Carcinoma	BLCA	blca_20may13_test	blca_tcga_pub
Brain Lower Grade Glioma	LGG	lgg_04oct13_seq	lgg_tcga
Breast invasive carcinoma	BRCA	brca_manuscript_rerun_nov12d_pw	brca_tcga_pub
Cervical squamous cell carcinoma and endocervical adenocarcinoma	CESC		cesc_tcga
Colon adenocarcinoma	COAD	coad_03feb13_seq_tumor_only	coadread_tcga
Esophageal carcinoma	ESCA		esca_tcga
Glioblastoma multiforme	GBM	gbm_2013_pub_tumor_only	gbm_tcga_pub2013
Head and Neck squamous cell carcinoma	HNSC	hnsc_03feb13_seq_tumor_only	hnsc_tcga
Kidney Chromophobe	KICH		kich_tcga_pub
Kidney renal clear cell carcinoma	KIRC	kirc_01oct12_A_pw	kirc_tcga_pub
Kidney renal papillary cell carcinoma	KIRP		kirp_tcga

Description	Name	Regulome Explorer	cBioPortal
Liver hepatocellular carcinoma	LIHC		lihc_tcga
Lung adenocarcinoma	LUAD	luad_03feb13_seq_tumor_only	luad_tcga_pub
Lung squamous cell carcinoma	LUSC	lusc_03feb13_seq_tumor_only	lusc_tcga
Ovarian serous cystadenocarcinoma	OV	ov_03feb13_ary_tumor_only	ov_tcga_pub
Pancreatic adenocarcinoma	PAAD		paad_tcga
Pheochromocytoma and Paraganglioma	PCPG		pcpg_tcga
Prostate adenocarcinoma	PRAD		prad_tcga
Rectum adenocarcinoma	READ		coadread_tcga
Sarcoma	SARC		sarc_tcga
Skin Cutaneous Melanoma	SKCM		skcm_tcga
Stomach adenocarcinoma	STAD	stad_23jan14_seq_tumor_only	stad_tcga
Thyroid carcinoma	THCA	thca_18oct14_TP	thca_tcga
Uterine Carcinosarcoma	UCS		ucs_tcga
Uterine Corpus Endometrial Carcinoma	UCEC	ucec_28jun13b_seq_tumor_only	ucec_tcga
Uveal Melanoma	UVM	uvm_20150515_private	

6. FREQUENTLY ASKED QUESTIONS

6.1. Wanderer says there are not enough samples to perform statistical analysis, what does it mean?

This means there were not enough number of informative samples to compute the Wilcoxon test. We note that, although we calculate the test with small number of samples, this result might be meaningless without a minimum number of cases.

6.2. The CSV files are a single line file!

We use Linux/UNIX linefeeds. Although your text viewer (i.e. notepad) might merge all the lines under Windows, your spreadsheet software (i.e. Excel) will properly recognize the line separation.

6.3. How can I open a CSV in my spreadsheet software?

In most cases it should work by just using the open with (right click with the mouse on the file icon). Alternatively, launch the application and import data from text, comma-separated or character-delimited

values (this depends on the software, i.e. LibreOffice).

6.4. How and when did you download the data from the TCGA?

Data from TCGA was downloaded using the TCGA-Assembler (Zhu et al., 2014) on July 8th 2014.

6.6. What is the origin of annotation data?

450k methylation array probes have been annotated as described in (Price et al., 2013).

Expression data annotation was performed using gene annotations and exon positions as provided by the TCGA and UCSC Genome Browser. Data was fetched using biomaRt package from R and, when needed, was merged according to feature overlaps (Gel et al., 2015).

6.6. What do RPKM and RSEM stand for?

Both values are directly obtained from the TCGA pipeline for RNA seq v2 expression analysis. RPKM stands for *reads per kilobase per million mapped reads* (Guo et al., 2013) and is used to represent exon expression levels. RSEM stands for *RNA-Seq by Expectation-Maximization* (Guo et al., 2013) and is used to represent gene expression levels. We log₂-transform these values adding one to get rid of zeroes; that means we plot log₂(x+1) RPKM or RSEM values.

7. REFERENCES

(Gel et al., 2015) Gel B., Diez-Villanueva A., Serra E., Buschbeck M., Peinado M.A. and Malinverni R. regioneR: an R/Bioconductor package for the association analysis of genomic regions based on permutation tests. Under review in Bioinformatics (2015)

(Guo et al., 2013) Guo Y., Sheng Q., Li J., Ye F., Samuels D.C. and Shyr Y. Large scale comparison of gene expression levels by microarrays and RNAseq using TCGA data. PloS One (2013), 8:8 doi:10.1371/journal.pone.0071462.

(Price et al., 2013) Price M.E., Cotton A.M., Lam L.L., FarrÃ© P., Emberly E., Brown C.J., Robinson W.P. and Kobor M.S. Additional annotation enhances potential for biologically-relevant analysis of the Illumina Infinium HumanMethylation450 BeadChip array. Epigenetics & Chromatin (2013), 6:4 doi:10.1186/1756-8935-6-4.

(Zeeberg et al., 2004) Zeeberg .BR., Riss J., Kane D.W., Bussey K.J., Uchio E., Linehan W.M., Barrett J.C. and Weinstein JN. Mistaken identifiers: gene name errors can be introduced inadvertently when using Excel in bioinformatics. BMC Bioinformatics (2004) Jun 23;5:80.

(Zhu et al., 2014) Zhu Y., Qiu P., Ji Y. TCGA-Assembler: Open-Source Software for Retrieving and Processing TCGA Data. Nature Methods (2014) 11:599-600 doi:10.1038/nmeth.2956.

8. VERSIONS AND COPYRIGHT

This documentation corresponds to Wanderer v1.0. The full development log can be found at <https://sourceforge.net/projects/tcga-wanderer/>. Wanderer is free software and is provided under the GPL v2 terms.

The previous major releases are:

- Wanderer v1.0 Wed Jun 10 10:56:31 2015 +0200 (commit 9339d9ee59d6fabaf802b1d17caaadb82786442)

Fixes minor bugs on profiling, allows data download in a ZIP file, adds new datasets, includes RNA seq by-gene data and correlates methylation and expression.

- Alpha Release Mon Dec 29 15:14:48 2014 (commit 21f1c7d9c742011e289842752a1ea871c13dc5fb)

Profiles DNA methylation and expression for matched normal-tumor datasets.

- Project start Mon Oct 6 15:04:45 2014 (commit 3e5440a552d04319331415e74635e144045cc6ae)

Project prototype.

This document corresponds to

```
commit 9339d9ee59d6fabaf802b1d17caaadb82786442 Author: Izaskun Mallona <imallona@imppc.org> Date:
Wed Jun 10 10:56:31 2015 +0200
```

Some datasets have limitations for usage until a global analysis is published; please contact TCGA before publishing.

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9. CONTACT

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We provide more tools and data at <http://www.maplab.cat>.